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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/781,928	02/20/2004	Connie Li Sun	034536-0829	9693
7590 11/25/2005			EXAMINER	
Stephen D. Pro	odnuk, Esq.	TUCKER, ZACHARY C		
Pfizer, Inc.			ART UNIT	DARED MIR (DED
Pfizer La Jolla Labs			ARTONII	PAPER NUMBER
10777 Science (1624		
San Diego, CA 92121			DATE MAILED: 11/25/2005	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)
	10/781,928	SUN ET AL.
Office Action Summary	Examiner	Art Unit
·		1624
The MAILING DATE of this communication ap	Zachary C. Tucker	
Period for Reply		
A SHORTENED STATUTORY PERIOD FOR REPI WHICHEVER IS LONGER, FROM THE MAILING [- Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. If NO period for reply is specified above, the maximum statutory period. Failure to reply within the set or extended period for reply will, by statu Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNIC .136(a). In no event, however, may a red will apply and will expire SIX (6) MONT te, cause the application to become ABA	ATION. ply be timely filed HS from the mailing date of this communication. INDONED (35 U.S.C. § 133).
Status		
3) Since this application is in condition for allowed	 is action is non-final. ance except for formal matte	• •
closed in accordance with the practice under	Ex parte Quayle, 1935 C.D.	11, 453 O.G. 213.
Disposition of Claims		
4) ⊠ Claim(s) 1-16 is/are pending in the application 4a) Of the above claim(s) is/are withdra 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 1-16 are subject to restriction and/or	awn from consideration.	
Application Papers		
9)⊠ The specification is objected to by the Examin	· ner	•
10) The drawing(s) filed on is/are: a) ac		y the Examiner.
Applicant may not request that any objection to the	•	` `
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the E	· · · · · · · · · · · · · · · · · · ·	
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority document application from the International Bureat * See the attached detailed Office action for a list	nts have been received. Its have been received in Appority documents have been reu au (PCT Rule 17.2(a)).	plication No eceived in this National Stage
Attachment(s)		
Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948)	4) 🔲 Interview Su Paper No(s)	mmary (PTO-413) Mail Date
 Notice of Draitsperson's Fatent Drawing Review (PTO-946) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08 Paper No(s)/Mail Date <u>1Sep04</u>. 		ormal Patent Application (PTO-152)

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- Claims 1-7, drawn to imidazopyrazines compounds, classified in class/subclass 544/350.
- II. Claims 8-16, drawn to methods of treating divers diseases and conditions, and a pharmaceutical composition employed for that purpose, classified in class/subclass 514/249.

The inventions are distinct, each from the other because:

Inventions I and II are related as product and process of use/ related sub-product not. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case conditions treated by methods in Group II are treatable by materially different means, such as various cancers being treated with cytotoxic agents. The sub-product of claim 8 has been included in Group II because of the similar classification of the pharmaceutical composition of that claim and the methods also included in Group II. Compounds according to Group I are not limited in their utility to only active ingredients in pharmaceutical compositions, as the paragraph bridging pages 42 and 42 of the instant specification (section [0126]), teaches.

So, a search required for simple disclosures of chemical compounds in Group I is not co-extensive with the search required for disclosures of methods and

pharmaceutical compositions according to Group II. A search required to determine compliance with the first paragraph of 35 U.S.C. 112, is not required for Group I. Such a search will entail a survey of the medical literature relating the use of certain kinase inhibitors in the treatment of disease, at the time the invention was made.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter and classification, restriction for examination purposes as indicated is proper.

This Requirement is further set forth as follows:

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Because all of substituents R₁ through R₆ are defined in expansive terms, the search required for claim 1, absent this additional Requirement for an Election of Species, would pose an undue burden on the examiner.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims

are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Upon election of a single disclosed species for examination, the search will proceed as described in MPEP 803.02, under "Markush Practice." The search will not be unnecessarily broadened.

Specification

The abstract of the disclosure is objected to because no generic structure of the compounds disclosed in the application is provided. As such, the abstract does not provide a concise description of the claimed and disclosed subject matter; the abstract is no more descriptive than the title of the application. Correction is required. See MPEP § 608.01(b).

Conclusion

Any inquiry concerning this communication should be directed to Zachary Tucker whose telephone number is (571) 272-0677. The examiner can normally be reached Tuesday-Thursday from 8:00am to 4:30pm or Monday from 6:00am to 1:30pm. If Attempts to reach the examiner are unsuccessful, contact the examiner's supervisor, James O. Wilson, at (571) 272-0661.

The fax number for the organization where this application or proceeding is assigned is (571) 273-8300.

Application/Control Number: 10/781,928

Art Unit: 1624

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600.

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